ORIGINAL RESEARCH

Laser Doppler Flowmetry and Visible Light Spectroscopy of the Gastric Tube During Minimally Invasive Esophagectomy

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Patients and Methods: Ten patients with esophageal cancer operated with minimally invasive esophagectomy participated in this single-center, prospective, observational pilot study. A single probe with laser Doppler flowmetry and visible light spectroscopy was used to perform transserosal microcirculation assessment of the gastric tube at predefined anatomical sites during different operation phases. Group comparison and changes were evaluated using the paired sample *t*-test.

Results: A reduction in StO2 was found at all measuring sites after the gastric tube formation compared with the baseline measurements. The mean StO2 reduction from baseline to gastric tube formation and after anastomosis was 16% (range 4%–28%) and 42% (range, 35%–52%), respectively. A statistically significant increase in the rHb concentration, representing venous congestion, was detected at the most cranial part of the gastric tube (*P* = 0.04). Three patients developed anastomotic leaks.

Conclusion: Intraoperative real-time laser Doppler flowmetry and visible light spectroscopy are feasible and may provide insight to microcirculatory changes in the gastric tube and at the anastomotic site. Patients with anastomotic leaks seem to have critical local tissue StO2 reduction and venous congestion that should be further evaluated in studies with larger sample sizes.

Keywords: esophagectomy, gastric tube circulation, gastroesophageal anastomosis complications

Introduction

Anastomotic leak is a severe complication after thoracic gastroesophageal anastomosis (TGEA) and is typically reported at incidences of 10–20%.^{1–3} Although multifactorial, ischemia at the anastomotic site plays a central role in TGEA leaks.⁴ The cranial 20% of the gastric tube is vascularized by an intramural vascular network and may be prone to the development of ischemia by manipulation, tension, and strangulation during gastric pull-up and anastomosis construction.^{3,5} The ischemic changes may not be visually apparent during the surgical procedure, and constructing the anastomosis at a hypoperfused site may result in an anastomotic leak. Intraoperative information about the gastric tube's microcirculatory status may guide the surgeon to define the

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anastomotic site with the best possible microcirculation. Despite the introduction of numerous intraoperative bowel viability assessment techniques, only a few techniques are feasible for intraoperative assessment of the gastric tube during esophagectomy.^{6,7} In addition to laser Doppler flowmetry (LDF), which is one of the methods most extensively investigated, visible light spectroscopy (VLS) has emerged as a promising method for the detection of the microcirculatory changes in the gastrointestinal tract.^{4,8–10}

The main aim of our pilot study was to evaluate the feasibility of combined use of LDF and VLS for transserosal microcirculation assessment of the stomach and the gastric tube during different phases of minimally invasive esophagectomy.

Patients and Methods

During seven months from July 2018 to January 2019, patients with esophageal cancer, scheduled for minimally invasive esophagectomy at Oslo University Hospital, Ullevål, were considered for this pilot study enrollment, at the discretion of the attending surgeon and from the availability of the principal investigator at the time of surgery. The institution is a regional center for esophageal cancer's surgical treatment, with an annual volume of about 50 patients operated. Inclusion criteria for study enrollment included patients with the capacity to give informed consent and a potentially curable distal esophageal cancer. All patients received neoadjuvant radio-chemotherapy with 41.1 Gy and Carboplatin and paclitaxel according to the CROSS radio-chemotherapy regime.¹¹

During surgery, a 2.6 mm microprobe with combined LDF and VLS modalities (O2C; LEA Medizintechnik, Germany), was channeled through a laparoscopic trocar to quantify transserosal, blood flow, velocity, mixed arterial and venous saturation of hemoglobin (StO2), and the amount of hemoglobin per tissue volume (rHb).

The O2C transmits continuous wave laser light (500– 630 nm) and white light (830 nm) through an optical fiber to the tissue, where it is scattered and collected with fibers of the probe placed on the tissue surface.¹² The white light tends to penetrate deeper into the tissue than the laser light due to its shorter wavelength.¹³ In VLS, the principle of absorbance and scattering of white light in biological tissues, gives a marked difference in absorption spectra of oxygenated and deoxygenated hemoglobin thereby, directly measures hemoglobin saturation and concentration.^{8,13,14}

For LDF measurements, reflected laser light from the moving red blood cells in the tissue generates a Doppler shift. The frequency of this reflected light is dependent upon the velocity of the cells (erythrocytes) and is detected by a photodetector within the instrument and transformed into an electrical signal. The LDF produces a value referred to as a flow (red blood cell flux) expressed as mL/min/100-gram tissue.

Measurements were repeated in case of unstable or fluctuating recordings. The graphical picture provided by the LCD monitor of the O2C unit, help to keep the absorption spectra of oxyhemoglobin well above 50% of the arbitrary unit (AU) scale during examinations (Figure 1A and B).¹⁵ This step allowed us to perform the repeated transserosal measurements in all subjects without applying unnecessary pressure on the serosa. Before each recording, an ambient light correction was performed automatically, which allowed keeping the examined area illuminated and maintaining the O2C microprobe's visual control throughout the examination. A standard measurement protocol of 5 seconds of continuous measurements at each anatomical position resulted in approximately 200 measurements at each



Figure I (A–B). Transserosal microcirculation recordings: (A) microprobe on ventricle surface (B) LCD monitor with the real-time absorption spectrum of oxyhemoglobin (red), and graphic presentation of StO2, relative hemoglobin, flow and velocity and the mean numerical values.

anatomical site. The system provided a real-time quantitative measurement and stored the raw data for later analysis.

After establishing pneumoperitoneum with CO2, and before any intraperitoneal dissection, baseline measurements were recorded from predefined anatomical sites on the greater curvature's anterior surface. Measurements were repeated at the same anatomical sites after gastric tube formation and subsequently, after the construction of the TGEA (Figure 2A–C). The distance between each measuring site was approximately 3–4 cm apart. In all patients, the gastroesophageal anastomosis was constructed at the site M7. A marking suture was placed at the gastric incisura towards the level of site M3, to identify the site after the gastric pull-up and anastomosis. All measurements were performed under stable hemodynamic conditions. No vasopressor medications were administered during the measurements, and the systemic oxygen saturation was kept > 97%.

Operative Technique

A standard thoracolaparoscopic, "Ivor-Lewis" type minimal invasive esophagectomy was performed in all patients.¹⁶ This approach included complete mobilization of the stomach, dissection of the short gastric arteries and the left gastric artery, and gastric tube preparation after regional lymphadenectomy. A multi-step thoracolaparoscopic subtotal resection of the esophagus with two-field lymphadenectomy was performed. The gastric conduit was anastomosed to the proximal residual esophagus at the carina level by a circular staple device. The introduction site for the circular stapler was closed using a linear stapler and oversewn.



Figure 2 (A-C). Measuring points M1 to M8 (green dots) at baseline, after gastric tube construction, and gastroesophageal anastomosis.

All surgical procedures were performed under general anesthesia with the same team of surgeons who were blinded to the results of the perioperative microcirculation measurements. Thoracic drains were placed close to the anastomosis and the diaphragmatic hiatus. A decompressing nasogastric tube was positioned, and the patients were extubated in the operating theatre. Postoperatively, a mean systemic arterial pressure of 65 mmHg or higher was targeted, and vasopressors were administered if needed. All patients stayed at the postoperative surveillance department for three days and were routinely examined on the third postoperative day with upper endoscopy and computed tomography (CT) with oral contrast of the esophagus. Complications are reported on and graded according to Clavien-Dindo classification of surgical complications as recommended the Esophagectomy Complications Consensus by Group.¹⁷

The study protocol was approved by the Regional Committees for Medical and Health Research Ethics in the South-Eastern region of Norway (approval number 2018/500/REK sør-øst A) and registered in Clinicaltrials. gov (ClinicalTrials.gov ID NCT03724162). The study conforms to the provisions of the Declaration of Helsinki. Informed written consent was obtained from all patients included in the study.

Statistical Analysis

A descriptive data analysis was performed. Data are presented as median (range) or mean (standard deviation) dependent on data distribution. Group comparison and mean changes were evaluated using the paired sample *t*-test. A *P* -value < 0.05 was considered statistically significant. Statistical analyses were performed using SPSS version 25 (IBM SPSS Statistics).

Results

During a period of seven months, ten patients with esophageal cancer were included. Patient characteristics are given in Table 1. There was no 30-day and 90-day mortality. During a median follow-up period of 16 months (range, 12–18 months), one patient died. Three patients (30%) had an anastomotic leak type I and Grade II surgical complication (Table 1). These three leakages were identified by standard upper endoscopy on the third postoperative day and confirmed with a CT with oral contrast on postoperative day six. In one patient, the leakage was treated solely with antibiotics. In the other two Table I Baseline Characteristics of Patients (n=10)

Variables						
Median age, years	59	59 (47–83)				
Gender (male: female)	3:7					
Comorbidity						
Ischemic heart disease	0					
Atrial fibrillation	Ι					
Stroke	0					
Hypertension	5					
Pulmonary disease		5				
Hypercholesterolemia	2	2				
BMI, median	24	.8 (2	1.6–31	.I)		
GERD	7	7				
Smoking	2					
Histology						
AC/SCC	8/2	8/2				
Tumor location (Distal Esophagus/Cardia)	9/1					
Clinical cancer staging						
Stage I	Т					
Stage II	3					
Stage III	5					
Stage IV	Т					
Type of operation						
Ivor-Lewis MIE	10	10				
Anastomotic leak						
Definition: Type I	3					
Clavien–Dindo classification of surgical	Ι	П	Illa	IIIb		
complications						
Anastomotic leak	0	3	0	0		
Pneumonia	0	3	0	0		
Pleural effusion	0	2	8	0		
Atrial fibrillation	0	4	0	0		
Pulmonary embolism	0	Т	0	0		
^a Median ICU stay, days	4 (4 (4–10)				
^b Median ICU stay, days		4 (4–12)				
^a Median hospital stay, days		25 (25–34)				
^b Median hospital stay, days		17 (7–39)				
Mortality						
30 days and 90 days	0					

Abbreviations: AC, adenocarcinoma; ASA, American Society of Anesthesiologists; BMI, body mass index; GERD, gastroesophageal reflux disease; ICU, intensive care unit; SCC, squamous cell carcinoma; ^aWith leaks; ^bWithout leaks.

patients, antibiotics were administered, and the Jackson-Prat drain kept for an extended period of 5–7 postoperative days. Intraoperative transserosal microcirculation assessment was successfully performed in all patients. The median recording time required for the measurements was 6 minutes (range, 3–8 minutes).

When analyzing the whole study group, a reduction in StO2 was found at all measuring sites in the gastric tube as compared with the baseline measurements. The reduction was statistically significant at the sites M4 (P = 0.04), M5, and M7 (P = 0.03). The reduction in StO2 became amplified (P < 0.001) after the construction of TGEA at all measuring sites as compared with the baseline measurements (Figure 3A). The mean StO2 reduction from baseline to gastric tube formation and after anastomosis was 16% (range 4% - 28%) and 42% (range, 35% - 52%), respectively. A statistically significant increase in the rHb concentration was detected at the most cranial part of the gastric tube, site M7 (P = 0.04) (Figure 3B).

The mean LDF measurements of the whole study group showed a statistically significant increase in the local blood flow at the site M4 (P = 0.009) after TGEA (Figure 3C). A similar change in velocity measurements was observed in the gastric tube after TGEA (P = 0.004) (Figure 3D).

In the three patients with leaks, the mean StO2 reduction after anastomosis, as compared with the baseline StO2, was 49% (range, 25% - 69%), while in patients without leaks, it was 39% (range, 32% - 46%). After anastomosis, rHb increased from baseline to 61% (range, 33% - 147%), and 17% (range, 0-38%), respectively in the patients with and without leaks. Although, the mean change in velocity was similar in patients with or without leaks, respectively 12% (range, -13% - 41%) and 11% (range, -14% - 44%), the mean tissue blood flow after anastomosis was increased by 36% (range, -5% - 57%) and 26% (range, -4% - 51%) respectively. This increase was statistically significant only at M4 in patients without leaks (P = 0.02). Table 2 gives a detailed account of microcirculation assessment results in patients with and without anastomotic leaks.

Discussion

This is the first study to present data on the simultaneous use of LDF and VLS for assessment of the gastric tube microcirculation in patients with esophageal cancer undergoing minimally invasive surgery. The O2C technology utilized, provides a real-time information to the surgeons



Figure 3 (A–D). Mean values with standard deviation of intraoperative transserosal microcirculation of stomach and gastric tube in patients with esophageal cancer. AU, arbitrary units; MI-M7, measuring sites.

Oxygen saturation (AU) (Mean ± SD)			^a Mean diff. in %	Р	P Relative Hemoglobin (AU) (Mean ± SD)			^a Mean diff. in %	Р	
	Baseline	Gastric tube	Anastomosis			Baseline	Gastric tube	Anastomosis	1	
MI ^b	69 ± 15	72 ± 27				80 ± 20	78 ± 13			
M۱۲	87 ± 9	78 ± 22				60 ± 15	63 ± 22			
M2 ^b	88 ± 7	76 ± 33				76 ± 14	58 ± 16			
M2 ^c	88 ± 6	69 ± 21				46 ± 14	65 ± 21			
M3 ^b	84 ± 7	90 ± 1	63 ± 28	25	.26	60 ± 16	81 ± 11	85 ± 35	42	.2
M3 ^c	90 ± 8	83 ± 8	54 ± 16	40	.005	62 ± 11	62 ± 11	62 ± 23	0	.2
M4 ^b	82 ± 12	74 ± 21	44 ± 39	46	.16	58 ± 15	77 ± 11	81 ± 21	40	.3
M4 ^c	91 ± 4	73 ± 16	51 ± 11	44	.005	48 ± 13	51 ± 20	55 ± 27	15	.5
M5 ^b	93 ± 2	67 ± 13	55 ± 3	41	.005	56 ± 16	69 ± 19	81 ± 10	46	.1
M5℃	89 ± 9	77 ± 17	60 ± 16	33	.005	45 ± 7	9 ± 19	58 ± 36	29	.4
M6 ^b	85 ± 12	81 ± 16	26 ± 12	69	.01	63 ± 31	62 ± 22	84 ± 6	33	.4
M6 ^c	86 ± 14	73 ± 16	59 ± 13	32	.005	48 ± 11	56 ± 17	49 ± 16	3	.8
M7 ^b	87 ± 7	85 ± 14	30 ± 14	66	.005	58 ± 24	47 ± 9	143 ± 77	15	.2
M7 ^c	90 ± 5	56 ± 29	49 ± 29	46	.01	40 ± 11	78 ± 23	55 ± 19	38	.1
Flow (AU) (Mean ± SD)					Velocity (AU) (Mean ± SD)					
				^a Mean diff. in %	Р		-		^a Mean diff. in %	Р
		Gastric tube	Anastomosis		Ρ		-	Anastomosis		Р
	n ± SD)	Gastric tube	Anastomosis		P	(Mean ± S	D)	Anastomosis		P
(Mea	n ± SD) Baseline		Anastomosis		P	(Mean ± S Baseline	D) Gastric tube	Anastomosis		P
(Mea MI ^b	Baseline 373 ± 139	290 ± 53	Anastomosis		P	(Mean ± S Baseline 34 ± 11	D) Gastric tube 34 ± 8	Anastomosis		P
(Mea MI ^b MI ^c	Baseline 373 ± 139 235 ± 69	290 ± 53 237 ± 70	Anastomosis		P	(Mean ± S Baseline 34 ± 11 34 ± 6	Gastric tube 34 ± 8 38 ± 19	Anastomosis		P
(Меа МІ ^ь МІ ^с М2 ^ь	Baseline 373 ± 139 235 ± 69 354 ± 90	290 ± 53 237 ± 70 313 ± 140	Anastomosis		P .9	(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14	Gastric tube 34 ± 8 38 ± 19 32 ± 2	Anastomosis 34 ± 6		Р .9
(Mea MI ^b MI ^c M2 ^b M2 ^c	n ± SD) Baseline 373 ± 139 235 ± 69 354 ± 90 249 ± 29	290 ± 53 237 ± 70 313 ± 140 252 ± 49		in %		(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14 35 ± 7	Gastric tube 34 ± 8 38 ± 19 32 ± 2 34 ± 5		in %	
(Mea MI ^b MI ^c M2 ^b M2 ^c M3 ^b	n ± SD) Baseline 373 ± 139 235 ± 69 354 ± 90 249 ± 29 307 ± 130	290 ± 53 237 ± 70 313 ± 140 252 ± 49 414 ± 251	292 ± 16	in %	.9	(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14 35 ± 7 32 ± 12	Gastric tube 34 ± 8 38 ± 19 32 ± 2 34 ± 5 50 ± 28	34 ± 6	in %	.9
(Mea MI ^b MI ^c M2 ^b M2 ^c M3 ^b M3 ^c	n ± SD) Baseline 373 ± 139 235 ± 69 354 ± 90 249 ± 29 307 ± 130 249 ± 65	$290 \pm 53 \\ 237 \pm 70 \\ 313 \pm 140 \\ 252 \pm 49 \\ 414 \pm 251 \\ 270 \pm 82$	292 ± 16 376 ± 183	in % -5 51	.9 .9	(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14 35 ± 7 32 ± 12 36 ± 12	Gastric tube 34 ± 8 38 ± 19 32 ± 2 34 ± 5 50 ± 28 38 ± 14	34 ± 6 46 ± 15	in % 5 28	.9 .005
(Mea MI ^b M2 ^b M2 ^c M3 ^b M3 ^c M4 ^b	n ± SD) Baseline 373 ± 139 235 ± 69 354 ± 90 249 ± 29 307 ± 130 249 ± 65 305 ± 70	$290 \pm 53 \\ 237 \pm 70 \\ 313 \pm 140 \\ 252 \pm 49 \\ 414 \pm 251 \\ 270 \pm 82 \\ 402 \pm 190 $	292 ± 16 376 ± 183 477 ± 135	in % -5 51 57	.9 .9 .3	(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14 35 ± 7 32 ± 12 36 ± 12 36 ± 14	Gastric tube 34 ± 8 38 ± 19 32 ± 2 34 ± 5 50 ± 28 38 ± 14 43 ± 12	34 ± 6 46 ± 15 45 ± 5	in % 5 28 25	.9 .005 .4
(Mea MI ^b M1 ^c M2 ^b M2 ^c M3 ^b M3 ^c M4 ^b M4 ^c	Baseline 373 ± 139 235 ± 69 354 ± 90 249 ± 29 307 ± 130 249 ± 65 305 ± 70 258 ± 41	$290 \pm 53 \\ 237 \pm 70 \\ 313 \pm 140 \\ 252 \pm 49 \\ 414 \pm 251 \\ 270 \pm 82 \\ 402 \pm 190 \\ 221 \pm 63$	292 ± 16 376 ± 183 477 ± 135 389 ± 117	-5 51 57 51	.9 .9 .3 .02	(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14 35 ± 7 32 ± 12 36 ± 12 36 ± 14 36 ± 4	Gastric tube 34 ± 8 38 ± 19 32 ± 2 34 ± 5 50 ± 28 38 ± 14 43 ± 12 30 ± 6	34 ± 6 46 ± 15 45 ± 5 52 ± 9	in % 5 28 25 47	.9 .005 .4 .005
(Mea MI ^b M2 ^b M2 ^c M3 ^b M3 ^c M4 ^b M4 ^c M5 ^b	n ± SD) Baseline 373 ± 139 235 ± 69 354 ± 90 249 ± 29 307 ± 130 249 ± 65 305 ± 70 258 ± 41 294 ± 80	$290 \pm 53 \\ 237 \pm 70 \\ 313 \pm 140 \\ 252 \pm 49 \\ 414 \pm 251 \\ 270 \pm 82 \\ 402 \pm 190 \\ 221 \pm 63 \\ 431 \pm 219$	292 ± 16 376 ± 183 477 ± 135 389 ± 117 427 ± 157	in % -5 51 57 51 45	.9 .9 .3 .02 .1	(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14 35 ± 7 32 ± 12 36 ± 12 36 ± 12 36 ± 14 36 ± 4 32 ± 3	Gastric tube 34 ± 8 38 ± 19 32 ± 2 34 ± 5 50 ± 28 38 ± 14 43 ± 12 30 ± 6 43 ± 12	34 ± 6 46 ± 15 45 ± 5 52 ± 9 45 ± 10	in % 5 28 25 47 39	.9 .005 .4 .005 .2
(Mea MI ^b MI ^c M2 ^b M2 ^c M3 ^b M3 ^c M4 ^b M4 ^c M5 ^b M5 ^c	Baseline 373 ± 139 235 ± 69 354 ± 90 249 ± 29 307 ± 130 249 ± 65 305 ± 70 258 ± 41 294 ± 80 227 ± 43	$290 \pm 53 \\ 237 \pm 70 \\ 313 \pm 140 \\ 252 \pm 49 \\ 414 \pm 251 \\ 270 \pm 82 \\ 402 \pm 190 \\ 221 \pm 63 \\ 431 \pm 219 \\ 249 \pm 66 \\ $	292 ± 16 376 ± 183 477 ± 135 389 ± 117 427 ± 157 262 ± 95	in % -5 51 57 51 45 15	.9 .9 .3 .02 .1 .4	(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14 35 ± 7 32 ± 12 36 ± 12 36 ± 14 36 ± 4 32 ± 3 34 ± 6	Gastric tube 34 ± 8 38 ± 19 32 ± 2 34 ± 5 50 ± 28 38 ± 14 43 ± 12 30 ± 6 43 ± 12 36 ± 7	$34 \pm 646 \pm 1545 \pm 552 \pm 945 \pm 1032 \pm 11$	in % 5 28 25 47 39 6	.9 .005 .4 .005 .2 .7
(Mea MI ^b M2 ^b M2 ^c M3 ^c M4 ^b M4 ^c M5 ^c M5 ^c	Baseline 373 ± 139 235 ± 69 354 ± 90 249 ± 29 307 ± 130 249 ± 65 305 ± 70 258 ± 41 294 ± 80 227 ± 43 263 ± 63	$290 \pm 53 \\ 237 \pm 70 \\ 313 \pm 140 \\ 252 \pm 49 \\ 414 \pm 251 \\ 270 \pm 82 \\ 402 \pm 190 \\ 221 \pm 63 \\ 431 \pm 219 \\ 249 \pm 66 \\ 282 \pm 173$	$292 \pm 16 \\ 376 \pm 183 \\ 477 \pm 135 \\ 389 \pm 117 \\ 427 \pm 157 \\ 262 \pm 95 \\ 346 \pm 152 $	in % -5 51 57 51 45 15 32	.9 .9 .3 .02 .1 .4 .5	(Mean ± S Baseline 34 ± 11 34 ± 6 44 ± 14 35 ± 7 32 ± 12 36 ± 12 36 ± 12 36 ± 4 32 ± 3 34 ± 6 31 ± 11	Gastric tube 34 ± 8 38 ± 19 32 ± 2 34 ± 5 50 ± 28 38 ± 14 43 ± 12 30 ± 6 43 ± 12 36 ± 7 31 ± 6	34 ± 6 46 ± 15 45 ± 5 52 ± 9 45 ± 10 32 ± 11 27 ± 3	in % 5 28 25 47 39 6 13	.9 .005 .4 .005 .2 .7 .6

Table 2 Results of Intraoperative Transserosal Microcirculation Assessment of Gastric Tube and Thoracic GastroesophagealAnastomosis with Laser Doppler Flowmetry and Visible Light Spectroscopy in Patients with Esophageal Cancer

Notes: P values, paired sample t test. ^aPercent changes comparison between Baseline and Anastomosis. ^bWith anastomotic leaks and ^cwithout leaks. **Abbreviation:** AU, arbitrary units.

during surgery. We found a significant reduction in the tissue StO2 in the gastric tube. The tissue StO2 deteriorated further, and a statistically significant reduction in the transversal StO2 after anastomosis was observed in all patients, compared to baseline values. In this pilot study, the patients with leaks had a lower mean StO2 at baseline compared with the patients without leaks, and they also had a more reduction in the mean StO2 after anastomosis.

The mean proportional increase in rHb in the patients with leaks was higher (61%) than those without leaks (17%). This increase in rHb was most evident in the most cranial part of the patients with leaks. This is an

important observation as the increase in rHb represents venous congestion that may impact tissue StO2. Gerau et al also found that in addition to significantly reduced StO2, there was an increase in the rHb in the gastric tube of patients with leaks.¹⁸ Furthermore, Buise et al found that the patients developed venous congestion after esophagectomy.¹⁹ Murakami et al performed microvascular anastomosis on the neck and showed that both the arterial circulation and venous congestion were relieved after such vascular anastomosis.^{20,21} However, in contradiction to Buise et al, they did not find reduced StO2. In the present study, we also found an increase in the mean

tissue blood flow from baseline to the anastomosis in patients with leaks (39%). This increase in tissue blood flow is of smaller magnitude in the patients without leaks (26%). Thus, the combined use of LDF and VLS enables substantiated information regarding blood circulation at the anastomotic site for the surgeon during surgery and may also further enlighten pathophysiological changes induced in the gastric tube.

Based on the present findings, patients with leaks may have more ischemic changes, aggravated by the venous congestion as represented by rHb increase. The increase in local blood flow is probably a compensatory physiological response to ischemia that may contribute to venous congestion due to reduced venous drainage caused by surgical trauma to the veins during esophagectomy and gastric tube formation. Extrinsic compression at the esophageal hiatus may also obstruct venous drainage and could have caused venous congestion in the patients with leaks.²¹ Manipulation and axial tension in the tube may also play a part in venous drainage.⁴ The physiological countercurrent mechanism of shunting blood flow from the mucosa to the serosal layer during an ischemic insult may also have contributed in the increase in local blood flow measured with transserosal LDF in our study cohort.²² To avoid vascular injury, a fair distance from the left gastroepiploic artery was maintained during free dissection of the greater omentum. Although the diaphragmatic hiatus's opening was wide, this could still have been a potential source of external compression on the venous drainage, secondary to expected postoperative edema.

The advantages of LDF and VLS are the quickness of measurements. low invasiveness. and high reproducibility.⁶ VLS is validated for the investigation of chronic mesenteric ischemia.¹⁴ Many studies have utilized LDF and VLS to assess the microcirculation in gastric tube. However, most of these studies of the microcirculatory assessment of gastric tube have been conducted on with the patients with anastomosis constructed on the neck. Most of the studies of the microcirculatory assessment of the gastric tubes are initial experiences and feasibility studies. Furthermore, either LDF or light spectrophotometry has been utilized. Only two studies incorporated both LDF and VLS.^{15,23} The former was the investigation of transmucosal microcirculation in patients with non-specific abdominal pain, and the latter included patients with chronic mesenteric ischemia and a control group with non-specific abdominal pain. Other studies such as Pham et al had no real-time measurements,

while Wang et al excluded patients with anastomotic leaks.^{24,25}

Although the assessment of ischemia can be identified as a common aim in these studies, the studies differ in patient demographics and the different measurement units. It is, therefore, hard to draw conclusions and to standardize these methods for routine clinical use. Interestingly, many of the available studies could not provide intraoperative real-time information about microcirculation.^{26,27} In contrast, the present study confirms the feasibility of the combined use of LDF and VLS intraoperatively, as made possible by the O2C technology.

Although studies have shown adverse effects of both radio- and chemotherapy on the tissue, the neoadjuvant radio-chemotherapy is the current standard of treatment of esophageal cancer before radical surgical resection for most patients.¹¹ Furthermore, the clinical results of major studies have so far not demonstrated more anastomotic leakages.²⁸ A major limitation of our pilot study is the small sample size. The study was not designed and powered to investigate anastomosis leaks in general. Therefore, statistical evaluations should be interpreted cautiously. Anastomotic leaks were identified early due to an aggressive diagnostic approach, including routine gastroscopy at day three and liberal use of early CT with oral contrast. The standardized surgical technique applied represents a strength of the study. The study results hold promise for future appropriately powered studies to provide intraoperative useful cut-off values for StO2, rHb, local tissue blood flow, and velocity. The results of more extensive studies may be used to standardize microcirculation assessment modalities for regular clinical use.

Conclusion

Concomitant intraoperative transserosal LDF and VLS may help identify local ischemia in the gastric tube during esophagectomy. Patients with anastomotic leaks seems to have a profound local tissue StO2 reduction, which is further aggravated by the development of venous congestion.

Data Sharing Statement

Individual participant data that underlie the results reported in this article, after deidentification (text, tables, figures), will be made available and shared with investigators whose proposed use of the data has been approved, but an independent review committee identified for this purpose. Proposals should be directed to associate professor Syed Sajid Hussain Kazmi MD Ph.D. sshkazmi@gmail.com, project leader. To gain access, data requesters will need to sign a data access agreement.

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Disclosure

The authors have nothing to disclose.

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